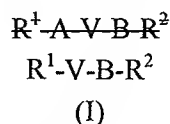


AMENDMENTS TO THE CLAIMS

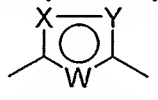
Listing of Claims:

This listing of claims will replace all prior versions and listings of the claims in the application.

1. (Currently Amended) A compound of formula (I), or a pharmaceutically acceptable salt thereof:



wherein V [[is]] represents a 5-membered heteroaryl ring containing up to four heteroatoms selected from O, N and S, optionally substituted by C₁₋₄ alkyl of the formula:



wherein W is N and one of X and Y is N and the other is O;

A is -CH=CH- or (CH₂)_n;

B is -CH=CH- or (CH₂)_n, where one of the CH₂ groups may be replaced by O, NR⁵, S(O)_m, C(O) or C(O)NR¹²;

n is ~~independently 0, 1, 2 or 3;~~

m is ~~independently 0, 1 or 2;~~

R¹ is ~~[[3- or]]~~4-pyridyl, ~~4- or 5-pyrimidinyl or 2-pyrazinyl, any of which may be optionally substituted by one or more substituents selected from 1 or 2~~ halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₃₋₇ cycloalkyl, aryl, OR⁶, CN, NO₂, S(O)_mR⁶, CON(R⁶)₂, N(R⁶)₂, NR¹⁰COR⁶, NR¹⁰SO₂R⁶, SO₂N(R⁶)₂, [[a]]4- to 7-membered heterocyclyl group or [[a]]5- or 6-membered heteroaryl groupgroups;

R² is 4- to 7-membered cycloalkyl substituted by R³, C(O)OR³, C(O)R³ or S(O)₂R³, or 4- to 7-membered heterocyclyl, containing one or two nitrogen atoms which is unsubstituted or substituted by C(O)OR⁴, C(O)R³, S(O)₂R³, C(O)NHR⁴, P(O)(OR¹¹)₂ or a 5- or 6-membered nitrogen containing heteroaryl group;

R³ is C₃₋₈ alkyl, C₃₋₈ alkenyl or C₃₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, heterocyclyl, heteroaryl, C₁₋₄ alkylC₃₋₇ cycloalkyl, C₁₋₄ alkylaryl, C₁₋₄ alkylheterocyclyl or C₁₋₄ alkylheteroaryl, any of which may be optionally substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶, CN, CO₂C₁₋₄ alkyl, N(R⁶)₂ and NO₂;

R⁴ is C₂₋₈ alkyl, C₂₋₈ alkenyl or C₂₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, heterocyclyl, heteroaryl, C₁₋₄ alkylC₃₋₇ cycloalkyl, C₁₋₄ alkylaryl, C₁₋₄ alkylheterocyclyl or C₁₋₄

alkylheteroaryl, any of which may be substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶, CN, CO₂C₁₋₄ alkyl, N(R⁶)₂ and NO₂;

R⁵ is hydrogen, C(O)R⁷, S(O)₂R⁸, C₃₋₇ cycloalkyl or C₁₋₄ alkyl optionally substituted by OR⁶, C₃₋₇ cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C₁₋₂ alkyl, C₁₋₂ fluoroalkyl, OR⁶, CN, N(R⁶)₂ and NO₂;

R⁶ are independently hydrogen C₁₋₄ alkyl, C₃₋₇ cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁹, CN, SO₂CH₃, N(R¹⁰)₂ and NO₂; or a group N(R¹⁰)₂ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR¹⁰;

R⁷ is hydrogen, C₁₋₄ alkyl, OR⁶, N(R⁶)₂, aryl or heteroaryl;

R⁸ is C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, aryl or heteroaryl;

R⁹ is hydrogen, C₁₋₂ alkyl or C₁₋₂ fluoroalkyl;

R¹⁰ is hydrogen or C₁₋₄ alkyl;

R¹¹ is phenyl; and

R¹² is hydrogen, C₁₋₄ alkyl or C₃₋₇ cycloalkyl;

provided that the compound is not:

- a) 4-(5-piperidin-4-yl-[1,2,4]oxadiazol-3-yl)pyridine;
- b) 4-(3-pyridin-4-yl-[1,2,4]oxadiazol-5-yl)piperidine-1-carboxylic acid ^tbutyl ester; or
- c) 4-[5-(4-butylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine[[:]]
- d) ~~3-[5-(4-butylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine; or~~
- e) ~~3-[5-(4-propylcyclohexyl)-[1,2,4]oxadiazol-3-yl]pyridine.~~

2-8. (Canceled)

9. (Currently Amended) A compound according to ~~claim 8~~claim 1, or a pharmaceutically acceptable salt thereof; wherein R¹ is 4-pyridyl optionally substituted by halo, C₁₋₄ alkyl, C₁₋₄ alkoxy or CN.

10. (Currently Amended) A compound according to ~~any one of the preceding claims~~claim 1, or a pharmaceutically acceptable salt thereof, wherein R² is a 4- to 7-membered cycloalkyl substituted by R³, or 4- to 7-membered heterocyclyl containing one nitrogen atom which is substituted by C(O)OR⁴.

11. (Currently Amended) A compound according to ~~any one of the preceding claims~~claim 1, or a pharmaceutically acceptable salt thereof, wherein R³ is C₃₋₈ alkyl which may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl.

12. (Currently Amended) A compound according to ~~any one of the preceding claims~~claim 1, or a pharmaceutically acceptable salt thereof, wherein R⁴ is C₂₋₈ alkyl, C₂₋₈ alkenyl or C₂₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, 5- to 6-membered heteroaryl containing one or two

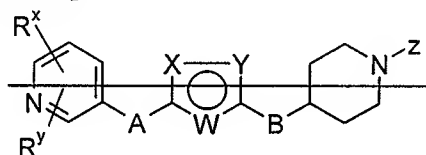
nitrogen atoms, C₁₋₄ alkyl, C₃₋₇ cycloalkyl or C₁₋₄ alkylaryl, any of which may be substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶ and CO₂C₁₋₄ alkyl.

13. (Original) A compound according to claim 12, or a pharmaceutically acceptable salt thereof, wherein R⁴ is C₃₋₆ alkyl optionally substituted with up to 5 fluoro or chloro atoms, and which may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl.

14. (Currently Amended) A compound according to ~~any one of the preceding claims~~ claim 1, or a pharmaceutically acceptable salt thereof, wherein R⁵ is C₁₋₄ alkyl.

15. (Currently Amended) A compound ~~of formula (I)~~ as defined in any one of Examples 1, ~~3 to 83 to 5, 10 to 13, 16 to 50~~ 16 to 39, 41, 42, or 52 to 149 ~~52 to 132, 134, 135, or 147 to 149~~, or a pharmaceutically acceptable salt thereof.

16. (Currently Amended) A compound according to claim 1 ~~having the formula (Id)~~, or a pharmaceutically acceptable salt thereof:



(Id)

where two of W, X and Y are N, and the other is O;

A is ~~CH=CH-~~ or (CH₂)_n;

wherein B is -CH=CH- or (CH₂)_n, where one of the CH₂ groups may be replaced by O, NR⁵, S(O)_m or C(O);

n is independently 0, 1, 2 or 3, ~~provided that not both n are 0~~;

m is independently 0, 1 or 2;

~~R^{*} and R^y are independently selected from hydrogen, halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₃₋₇ cycloalkyl, aryl, OR⁶, CN, NO₂, S(O)_mR⁶, CON(R⁶)₂, N(R⁶)₂, NR¹⁰COR⁶, NR¹⁰SO₂R⁶, SO₂N(R⁶)₂, a 4- to 7-membered heterocyclyl group and a 5- or 6-membered heteroaryl group;~~

R² is a 4- to 7-membered heterocyclyl containing one nitrogen atom which is substituted by C(O)OR⁴Z is C(O)OR⁴, C(O)R³, S(O)₂R³, C(O)NHR⁴ or a 5- or 6-membered nitrogen containing heteroaryl group;

~~R³ is C₃₋₈ alkyl, C₃₋₈ alkenyl or C₃₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, heterocyclyl, heteroaryl, C₁₋₄ alkyl, C₃₋₇ cycloalkyl, C₁₋₄ alkylaryl, C₁₋₄ alkylheterocyclyl or C₁₋₄ alkylheteroaryl, any of which may be optionally substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶, CN, CO₂C₁₋₄ alkyl, N(R⁶)₂ and NO₂;~~

R^4 is C_{2-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH_2 group that may be replaced by O, or C_{3-7} cycloalkyl, aryl, heterocyclyl, heteroaryl, C_{1-4} alkyl C_{3-7} cycloalkyl, C_{1-4} alkylaryl, C_{1-4} alkylheterocyclyl or C_{1-4} alkylheteroaryl, any of which may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^6 , CN, CO_2C_{1-4} alkyl, $N(R^6)_2$ and NO_2 ;

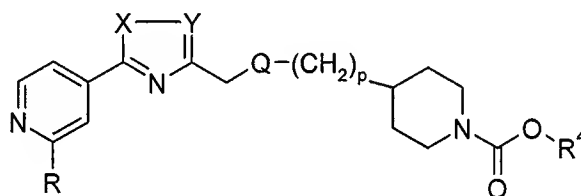
R^5 is hydrogen or C_{1-4} alkyl;

R^6 are independently hydrogen, or C_{1-4} alkyl, C_{3-7} cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^9 , CN, SO_2CH_3 , $N(R^{10})_2$ and NO_2 ; or a group $N(R^{10})_2$ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR^{10} ;

R^9 is hydrogen, C_{1-2} alkyl or C_{1-2} fluoroalkyl; and

R^{10} is hydrogen or C_{1-4} alkyl.

17. (Original) A compound according to claim 1 having the formula (Ie), or a pharmaceutically acceptable salt thereof:



(Ie)

wherein one of X and Y is N, and the other is O;

Q is O, NR^5 or CH_2 ;

R is hydrogen, halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{3-7} cycloalkyl, aryl, OR^6 , CN, NO_2 , $S(O)_mR^6$, $CON(R^6)_2$, $N(R^6)_2$, $NR^{10}COR^6$, $NR^{10}SO_2R^6$, $SO_2N(R^6)_2$, a 4- to 7-membered heterocyclyl group or a 5- or 6-membered heteroaryl group;

R^4 is C_{2-8} alkyl, C_{2-8} alkenyl or C_{2-8} alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and contain a CH_2 group that may be replaced by O, or C_{3-7} cycloalkyl, aryl, heterocyclyl, heteroaryl, C_{1-4} alkyl C_{3-7} cycloalkyl, C_{1-4} alkylaryl, C_{1-4} alkylheterocyclyl or C_{1-4} alkylheteroaryl, any of which may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^6 , CN, CO_2C_{1-4} alkyl, $N(R^6)_2$ and NO_2 ;

R^5 is C_{1-4} alkyl;

R^6 are independently hydrogen, or C_{1-4} alkyl, C_{3-7} cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C_{1-4} alkyl, C_{1-4} fluoroalkyl, OR^9 , CN, SO_2CH_3 , $N(R^{10})_2$ and NO_2 ; or a group $N(R^{10})_2$ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR^{10} ;

R^9 is hydrogen, C_{1-2} alkyl or C_{1-2} fluoroalkyl;

R^{10} is hydrogen or C_{1-4} alkyl; and

p is 0 or 1.

18. (Currently Amended) A pharmaceutical composition comprising a compound according to ~~any one of claims 1 to 17~~claim 1, including the ~~compounds~~compound of ~~provisos~~proviso c) to e), or a pharmaceutically acceptable salt thereof; and a pharmaceutically acceptable carrier.

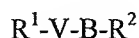
19. (Currently Amended) A method for the treatment of a disease or condition in which GPR116 plays a role comprising a step of administering to a subject in need thereof an effective amount of a compound according to ~~any one of claims 1 to 17~~claim 1, including the compounds of ~~provisos a) to e)~~a) to c), or a pharmaceutically acceptable salt thereof.

20. (Currently Amended) A method for the regulation of satiety comprising a step of administering to a subject in need thereof an effective amount of a compound according to ~~any one of claims 1 to 17~~claim 1, including the compounds of ~~provisos a) to e)~~a) to c), or a pharmaceutically acceptable salt thereof.

21. (Currently Amended) A method for the treatment of obesity comprising a step of administering to a subject in need thereof an effective amount of a compound according to ~~any one of claims 1 to 17~~claim 1, including the compounds of ~~provisos a) to e)~~a) to c), or a pharmaceutically acceptable salt thereof.

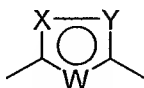
22. (Currently Amended) A method for the treatment of diabetes comprising a step of administering to a subject in need thereof an effective amount of a compound according to ~~any one of claims 1 to 17~~claim 1, including the compounds of ~~provisos a) to e)~~a) to c), or a pharmaceutically acceptable salt thereof.

23. (New) A method for the treatment of a disease or condition in which GPR116 plays a role comprising a step of administering to a subject in need thereof an effective amount of a compound of the formula:



or a pharmaceutically acceptable salt thereof;

wherein V represents a 5-membered heteroaryl ring of the formula:



wherein W is N and one of X and Y is N and the other is O;

B is $-\text{CH}=\text{CH}-$ or $(\text{CH}_2)_n$, where one of the CH_2 groups may be replaced by O, NR^5 , $\text{S}(\text{O})_m$, $\text{C}(\text{O})$ or $\text{C}(\text{O})\text{NR}^{12}$;

n is 0, 1, 2 or 3;

m is 0, 1 or 2;

R¹ is 3- or 4-pyridyl, 4- or 5-pyrimidinyl or 2-pyrazinyl, any of which may be optionally substituted by one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₃₋₇ cycloalkyl, aryl, OR⁶, CN, NO₂, S(O)_mR⁶, CON(R⁶)₂, N(R⁶)₂, NR¹⁰COR⁶, NR¹⁰SO₂R⁶, SO₂N(R⁶)₂, a 4- to 7-membered heterocyclyl group or a 5- or 6-membered heteroaryl group;

R² is 4- to 7-membered cycloalkyl substituted by R³, C(O)OR³, C(O)R³ or S(O)₂R³, or 4- to 7-membered heterocyclyl, containing one or two nitrogen atoms which is unsubstituted or substituted by C(O)OR⁴, C(O)R³, S(O)₂R³, C(O)NHR⁴, P(O)(OR¹¹)₂ or a 5- or 6-membered nitrogen containing heteroaryl group;

R³ is C₃₋₈ alkyl, C₃₋₈ alkenyl or C₃₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, heterocyclyl, heteroaryl, C₁₋₄ alkylC₃₋₇ cycloalkyl, C₁₋₄ alkylaryl, C₁₋₄ alkylheterocyclyl or C₁₋₄ alkylheteroaryl, any of which may be optionally substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶, CN, CO₂C₁₋₄ alkyl, N(R⁶)₂ and NO₂;

R⁴ is C₂₋₈ alkyl, C₂₋₈ alkenyl or C₂₋₈ alkynyl, any of which may be optionally substituted with up to 5 fluoro or chloro atoms, and may contain a CH₂ group that may be replaced by O, or C₃₋₇ cycloalkyl, aryl, heterocyclyl, heteroaryl, C₁₋₄ alkylC₃₋₇ cycloalkyl, C₁₋₄ alkylaryl, C₁₋₄ alkylheterocyclyl or C₁₋₄ alkylheteroaryl, any of which may be substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁶, CN, CO₂C₁₋₄ alkyl, N(R⁶)₂ and NO₂;

R⁵ is hydrogen, C(O)R⁷, S(O)₂R⁸, C₃₋₇ cycloalkyl or C₁₋₄ alkyl optionally substituted by OR⁶, C₃₋₇ cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C₁₋₂ alkyl, C₁₋₂ fluoroalkyl, OR⁶, CN, N(R⁶)₂ and NO₂;

R⁶ are independently hydrogen C₁₋₄ alkyl, C₃₋₇ cycloalkyl, aryl, heterocyclyl or heteroaryl, wherein the cyclic groups may be substituted with one or more substituents selected from halo, C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, OR⁹, CN, SO₂CH₃, N(R¹⁰)₂ and NO₂; or a group N(R¹⁰)₂ may form a 4- to 7-membered heterocyclic ring optionally containing a further heteroatom selected from O and NR¹⁰;

R⁷ is hydrogen, C₁₋₄ alkyl, OR⁶, N(R⁶)₂, aryl or heteroaryl;

R⁸ is C₁₋₄ alkyl, C₁₋₄ fluoroalkyl, aryl or heteroaryl;

R⁹ is hydrogen, C₁₋₂ alkyl or C₁₋₂ fluoroalkyl;

R¹⁰ is hydrogen or C₁₋₄ alkyl;

R¹¹ is phenyl; and

R¹² is hydrogen, C₁₋₄ alkyl or C₃₋₇ cycloalkyl.